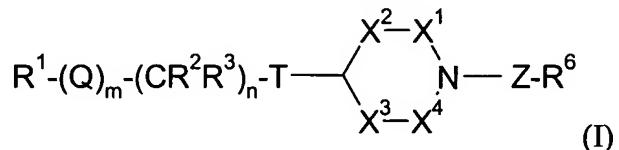


Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

1. (Presently amended) The present invention provides a A compound of formula (I), or a pharmaceutically acceptable salt thereof, or solvate thereof, or a solvate of a salt thereof:



wherein

Z is CR⁴R⁵, C(O) or CR⁴R⁵Z⁺, wherein R⁴ and R⁵ are CH₂;

Z⁺ is C₁₋₄ alkylene, C₂₋₄ alkenylene or C(O)NH;

R¹ represents a C₁-C₁₂ alkyl group optionally substituted by one or more substituents independently selected from cyano, hydroxyl, C₁-C₆ alkoxy, C₁-C₆ alkylthio, C₃₋₇ cycloalkyl, C₁-C₆ alkoxy carbonyl and phenyl (itself optionally substituted by one or more of halogen, nitro, cyano, C₁-C₆ alkyl, C₁-C₆ haloalkyl, phenyl(C₁-C₆ alkyl), C₁-C₆ alkoxy, C₁-C₆ haloalkoxy, S(O)₂(C₁-C₆ alkyl), C(O)NH₂, carboxy or C₁-C₆ alkoxy carbonyl); or

R¹ represents C₂-C₆ alkenyl optionally substituted by phenyl (itself optionally substituted by one or more of halogen, nitro, cyano, C₁-C₆ alkyl, C₁-C₆ haloalkyl, phenyl(C₁-C₆ alkyl), C₁-C₆ alkoxy, C₁-C₆ haloalkoxy, S(O)₂(C₁-C₆ alkyl), C(O)NH₂, carboxy or C₁-C₆ alkoxy carbonyl); or

R¹ represents a 3- to 14-membered saturated or unsaturated ring system which optionally comprises up to two ring carbon atoms that form carbonyl groups and which optionally further comprises up to 4 ring heteroatoms independently selected from nitrogen, oxygen and sulphur, wherein the ring system is optionally substituted by one or more substituents independently selected from: halogen, cyano, nitro, oxo, hydroxyl, C₁-C₈ alkyl, C₁-C₆ hydroxyalkyl, C₁-C₆

haloalkyl, C₁₋₆ alkoxy(C_{1-C₆} alkyl), C_{3-C₇} cycloalkyl(C_{1-C₆} alkyl), C_{1-C₆} alkylthio(C_{1-C₆} alkyl), C_{1-C₆} alkylcarbonyloxy(C_{1-C₆} alkyl), C_{1-C₆} alkylS(O)₂(C_{1-C₆} alkyl), aryl(C_{1-C₆} alkyl), heterocyclyl(C_{1-C₆} alkyl), arylS(O)₂(C_{1-C₆} alkyl), heterocyclylS(O)₂(C_{1-C₆} alkyl), aryl(C_{1-C₆} alkyl)S(O)₂, heterocyclyl(C_{1-C₆} alkyl)S(O)₂, C_{2-C₆} alkenyl, C_{1-C₆} alkoxy, carboxy-substituted C_{1-C₆} alkoxy, C_{1-C₆} haloalkoxy, C_{1-C₆} hydroxyalkoxy, C_{1-C₆} alkylcarboxy-substituted C_{1-C₆} alkoxy, aryloxy, heterocycloloxy, C_{1-C₆} alkylthio, C_{3-C₇} cycloalkyl(C_{1-C₆} alkylthio), C_{3-C₆} alkynylthio, C_{1-C₆} alkylcarbonylamino, C_{1-C₆} haloalkylcarbonylamino, SO₃H, -NR⁷R⁸, -C(O)NR²³R²⁴, S(O)₂NR¹⁸R¹⁹, S(O)₂R²⁰, R²⁵C(O), carboxyl, C_{1-C₆} alkoxycarbonyl, aryl and heterocyclyl; wherein the foregoing aryl and heterocyclyl moieties are optionally substituted by one or more of halogen, oxo, hydroxy, nitro, cyano, C_{1-C₆} alkyl, C_{1-C₆} haloalkyl, phenyl(C_{1-C₆} alkyl), C_{1-C₆} alkoxy, C_{1-C₆} haloalkoxy, S(O)₂(C_{1-C₆} alkyl), C(O)NH₂, carboxy or C_{1-C₆} alkoxycarbonyl;

m is 0 or 1;

~~Q represents an oxygen or sulphur atom or a group NR⁹, C(O), C(O)NR⁹, NR⁹C(O) or CH-CH;~~
n is 0, 1, 2, 3, 4, 5 or 6 provided that when n is 0, then m is 0;

each R² and R³ independently represents a hydrogen atom or a C_{1-C₄} alkyl group, or (CR²R³)_n represents C_{3-C₇} cycloalkyl optionally substituted by C_{1-C₄} alkyl;

T represents a group NR¹⁰, C(O)NR¹⁰, NR¹¹C(O)NR¹⁰ or C(O)NR¹⁰NR¹¹, wherein R¹⁰ is H; X¹, X², X³ and X⁴ are, independently, CH₂-CHR¹² (wherein each R¹² is, independently, C_{1-C₄} alkyl or C_{3-C₇} cycloalkyl(C_{1-C₄} alkyl)) or C=O; or, when they are CHR¹², the R¹² groups of X¹ and X³ or X⁴, or X² and X³ or X⁴ join to form a two or three atom chain which is CH₂CH₂, CH₂CH₂CH₂, CH₂OCH₂ or CH₂SCH₂; provided always that at least two of X¹, X², X³ and X⁴ are CH₂;

R⁴ and R⁵ each independently represent a hydrogen atom or a C_{1-C₄} alkyl group;

R⁶ is ~~aryl or heterocyclic phenyl~~, both optionally substituted by one or more of: halogen, cyano, nitro, oxo, hydroxyl, C_{1-C₈} alkyl, C_{1-C₆} hydroxyalkyl, C_{1-C₆} haloalkyl, C₁₋₆ alkoxy(C_{1-C₆} alkyl), C_{3-C₇} cycloalkyl(C_{1-C₆} alkyl), C_{1-C₆} alkylthio(C_{1-C₆} alkyl), C_{1-C₆} alkylcarbonyloxy(C_{1-C₆} alkyl), C_{1-C₆} alkylS(O)₂(C_{1-C₆} alkyl), aryl(C_{1-C₆} alkyl), heterocyclyl(C_{1-C₆} alkyl), arylS(O)₂(C_{1-C₆} alkyl);

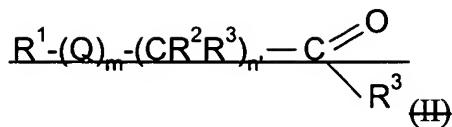
C_6 alkyl), heterocyclylS(O)₂(C_1 - C_6 alkyl), aryl(C_1 - C_6 alkyl)S(O)₂, heterocyclyl(C_1 - C_6 alkyl)S(O)₂, C_2 - C_6 alkenyl, C_1 - C_6 alkoxy, carboxy-substituted C_1 - C_6 alkoxy, C_1 - C_6 haloalkoxy, C_1 - C_6 hydroxyalkoxy, C_1 - C_6 alkylcarboxy-substituted C_1 - C_6 alkoxy, aryloxy, heterocyclyloxy, C_1 - C_6 alkylthio, C_3 - C_7 cycloalkyl(C_1 - C_6 alkylthio), C_3 - C_6 alkynylthio, C_1 - C_6 alkylcarbonylamino, C_1 - C_6 haloalkylcarbonylamino, SO₃H, -NR¹⁶R¹⁷, -C(O)NR²¹R²², S(O)₂NR¹³R¹⁴, S(O)₂R¹⁵, R²⁶C(O), carboxyl, C_1 - C_6 alkoxy carbonyl, aryl and heterocyclyl; wherein the foregoing aryl and heterocyclyl moieties are optionally substituted by one or more of halogen, nitro, cyano, C_1 - C_6 alkyl, C_1 - C_6 haloalkyl, phenyl(C_1 - C_6 alkyl), C_1 - C_6 alkoxy, C_1 - C_6 haloalkoxy, S(O)₂(C_1 - C_6 alkyl), C(O)NH₂, carboxy or C_1 - C_6 alkoxy carbonyl; R⁷, R⁸, R⁹, R¹⁰, R¹¹, R¹³, R¹⁴, R¹⁶, R¹⁷, R¹⁸, R¹⁹, R²¹, R²², R²³ and R²⁴ are, independently hydrogen, C_1 - C_6 alkyl, C_1 - C_6 haloalkyl, C_1 - C_6 hydroxyalkyl, C_3 - C_7 cycloalkyl, C_3 - C_7 cycloalkyl(C_1 - C_4 alkyl) or phenyl(C_1 - C_6 alkyl); and, R¹⁵ and R²⁰ are, independently, C_1 - C_6 alkyl, C_1 - C_6 hydroxyalkyl, C_3 - C_6 cycloalkyl, C_3 - C_7 cycloalkyl(C_1 - C_4 alkyl) or C_1 - C_6 alkyl optionally substituted by phenyl; R²⁵ and R²⁶ are, independently, C_1 - C_6 alkyl or phenyl (optionally substituted by one or more of halogen, nitro, cyano, C_1 - C_6 alkyl, C_1 - C_6 haloalkyl, phenyl(C_1 - C_6 alkyl), C_1 - C_6 alkoxy, C_1 - C_6 haloalkoxy, S(O)₂(C_1 - C_6 alkyl), C(O)NH₂, carboxy or C_1 - C_6 alkoxy carbonyl); or a pharmaceutically acceptable salt thereof, or solvate thereof, or a solvate of a salt thereof; provided that when T is C(O)NR¹⁰ and R⁺ is optionally substituted phenyl then n is not 0.

2-4. (Cancelled)

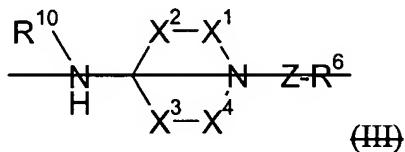
5. A compound as defined in any one of Examples 1 to 416.

6. (Presently amended) A process for the preparation of a compound of formula (I) as defined in claim 1 which comprises:

(a) when n is at least 1, the CR²R³ group attached directly to T is CHR³ and T is NR¹⁰, reacting a compound of general formula

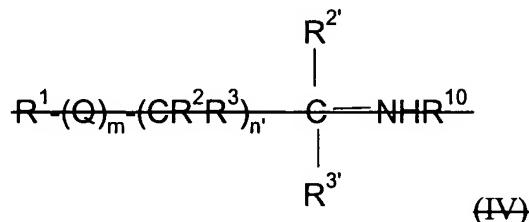


wherein n' is 0 or an integer from 1 to 3 and R^1 , R^2 , R^3 , m and Q are as defined in formula (I), with a compound of general formula

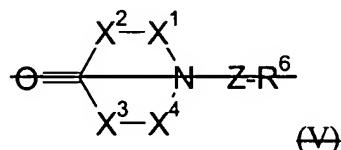


or a salt thereof, wherein X^1 , X^2 , X^3 , X^4 , Z , R^6 and R^{10} are as defined in formula (I), in the presence of a reducing agent; or

(b) when n is at least 1, the CR^2R^3 group attached directly to T is $\text{C}(\text{C}_1\text{-C}_4\text{ alkyl})_2$ and T is NR^{10} , reacting a compound of general formula

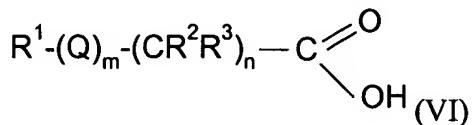


wherein n' is 0 or an integer from 1 to 3, $\text{R}^{2'}$ and $\text{R}^{3'}$ each independently represent a $\text{C}_1\text{-C}_4\text{ alkyl}$ group, and R^1 , R^2 , R^3 , R^{10} , m and Q are as defined in formula (I), with a compound of general formula

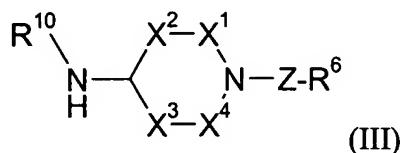


wherein X^1 , X^2 , X^3 , X^4 , Z and R^6 are as defined in formula (I), in the presence of a reducing agent; or

(ea) when T is $C(O)NR^{10}$, reacting a compound of general formula

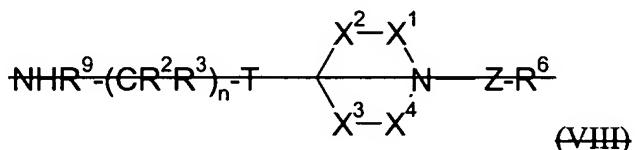


wherein R^1 , R^2 , R^3 , Q , m and n are as defined in formula (I), with a compound of formula (III)



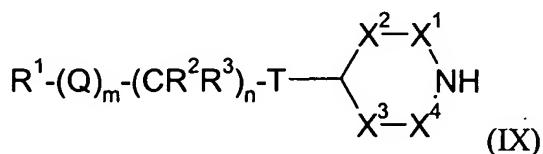
wherein X^1 , X^2 , X^3 , X^4 , Z , R^6 and R^{10} are as defined in formula (I),
or a salt thereof as defined in (a) above; or

(d) when m is 1 and Q is NR^9 , reacting a compound of general formula (VII), R^+-L^+ ,
wherein L^+ represents a leaving group (e.g. a halogen atom) and R^+ is as defined in formula (I),
with a compound of general formula



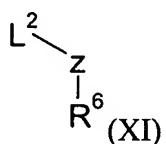
or a salt thereof, wherein n , T , X^1 , X^2 , X^3 , X^4 , Z , R^2 , R^3 , R^6 and R^9 are as defined in formula (I);
or

(eb) when at least one of R^4 and R^5 represents a hydrogen atom, reacting a compound of
general formula



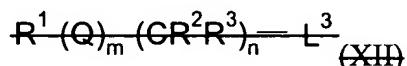
or a salt thereof, wherein R^1 , R^2 , R^3 , Q , m , n , X^1 , X^2 , X^3 , X^4 and T are as defined in formula (I), with a compound of general formula (X), $R^6 - C(O) - R^{20}$, wherein R^{20} represents a hydrogen atom or a C_1 - C_4 alkyl group and R^6 is as defined in formula (I), in the presence of a reducing agent; or

(fc) reacting a compound of formula (IX) as defined in (e)(b) above, with a compound of general formula



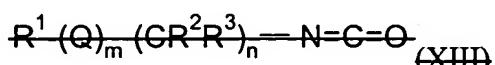
wherein L^2 represents a leaving group (e.g. a halogen atom) and Z and R^6 are as defined in formula (I); or

(g) when T is NR^{10} , reacting a compound of general formula



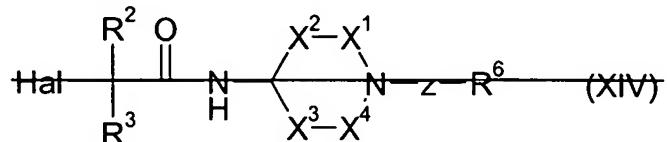
wherein L^3 represents a leaving group (e.g. a halogen atom) and R^1 , R^2 , R^3 , m , n and Q are as defined in formula (I), with a compound of formula (III) or a salt thereof as defined in (a) above; or

(h) when T is $NHC(O)NR^{10}$, reacting a compound of general formula



wherein R^1 , R^2 , R^3 , Q , m and n are as defined in formula (I), with a compound of formula (III) or a salt thereof as defined in (a) above; or

(+) ~~when T is C(O)NH, Z is CH₂, n is 1, R² and R³ are hydrogen or C₁-C₄ alkyl and Q is oxygen or sulphur, reacting a compound of formula (XIV):~~



~~wherein Hal is a suitable halogen, R², R³, X¹, X², X³, X⁴, Z and R⁶ are as defined in formula (I), with R¹OH or R¹SH in the presence of a suitable base;~~

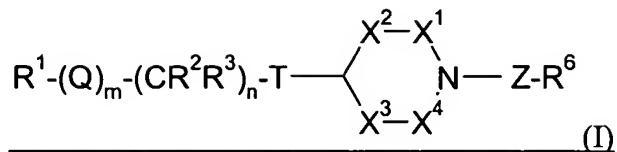
and optionally after (a), (b), or (c), ~~(d), (e), (f), (g), (h) or (i)~~ forming a pharmaceutically acceptable salt or solvate of the compound of formula (I) obtained.

7. (Presently amended) A pharmaceutical composition comprising a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as ~~claimed in any one of claims~~ claim 1 to 4 in association with a pharmaceutically acceptable adjuvant, diluent or carrier.

8. (Presently amended) A process for the preparation of a pharmaceutical composition as claimed in claim 7 which comprises mixing a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as ~~claimed in any one of claims~~ claim 1 to 4 with a pharmaceutically acceptable adjuvant, diluent or carrier.

9-10. (Cancelled)

11. (Presently Amended) A method of treating an inflammatory disease ~~asthma~~ in a patient suffering from, or at risk of, said disease, which comprises administering to the patient a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt thereof, or solvate thereof, or a solvate of a salt thereof, ~~as defined in claim 10.~~



wherein

Z is CR⁴R⁵, wherein R⁴ and R⁵ are CH₂;

R¹ represents a 3- to 14-membered saturated or unsaturated ring system which comprises up to two ring carbon atoms that form carbonyl groups and which further comprises up to 4 ring heteroatoms independently selected from nitrogen, oxygen and sulphur, wherein the ring system is optionally substituted by one or more substituents independently selected from: halogen, cyano, nitro, oxo, hydroxyl, C₁-C₈ alkyl, C₁-C₆ hydroxylalkyl, C₁-C₆ haloalkyl, C₁-C₆ alkoxy(C₁-C₆ alkyl), C₃-C₇ cycloalkyl(C₁-C₆ alkyl), C₁-C₆ alkylthio(C₁-C₆ alkyl), C₁-C₆ alkylcarbonyloxy(C₁-C₆ alkyl), C₁-C₆ alkylS(O)₂(C₁-C₆ alkyl), aryl(C₁-C₆ alkyl), heterocyclyl(C₁-C₆ alkyl), arylS(O)₂(C₁-C₆ alkyl), heterocyclylS(O)₂(C₁-C₆ alkyl), aryl(C₁-C₆ alkyl)S(O)₂, heterocyclyl(C₁-C₆ alkyl)S(O)₂, C₂-C₆ alkenyl, C₁-C₆ alkoxy, carboxy-substituted C₁-C₆ alkoxy, C₁-C₆ haloalkoxy, C₁-C₆ hydroxylalkoxy, C₁-C₆ alkylcarboxy-substituted C₁-C₆ alkoxy, aryloxy, heterocycloloxy, C₁-C₆ alkylthio, C₃-C₇ cycloalkyl(C₁-C₆ alkylthio), C₃-C₆ alkynylthio, C₁-C₆ alkylcarbonylamino, C₁-C₆ haloalkylcarbonylamino, SO₃H, -NR⁷R⁸, -C(O)NR²³R²⁴, S(O)₂NR¹⁸R¹⁹, S(O)₂R²⁰, R²⁵C(O), carboxyl, C₁-C₆ alkoxy carbonyl, aryl and heterocyclyl;
wherein the foregoing aryl and heterocyclyl moieties are optionally substituted by one or more of halogen, oxo, hydroxy, nitro, cyano, C₁-C₆ alkyl, C₁-C₆ haloalkyl, phenyl(C₁-C₆ alkyl), C₁-C₆ alkoxy, C₁-C₆ haloalkoxy, S(O)₂(C₁-C₆ alkyl), C(O)NH₂, carboxy or C₁-C₆ alkoxy carbonyl;

m is 0;

n is 2;

each R² and R³ independently represents a hydrogen atom or a C₁-C₄ alkyl group, or (CR²R³)_n represents C₃-C₇ cycloalkyl optionally substituted by C₁-C₄ alkyl;

T represents a group C(O)NR¹⁰;

X¹, X², X³ and X⁴ are, independently, CH₂;

R⁴ and R⁵ each independently represent a hydrogen atom or a C₁-C₄ alkyl group;

R⁶ is phenyl optionally substituted by one or more of: halogen, cyano, nitro, oxo, hydroxyl, C₁-C₈ alkyl, C₁-C₆ hydroxyalkyl, C₁-C₆ haloalkyl, C₁-C₆ alkoxy(C₁-C₆ alkyl), C₃-C₇ cycloalkyl(C₁-C₆ alkyl), C₁-C₆ alkylthio(C₁-C₆ alkyl), C₁-C₆ alkylcarbonyloxy(C₁-C₆ alkyl), C₁-C₆ alkylS(O)₂(C₁-C₆ alkyl), aryl(C₁-C₆ alkyl), heterocyclyl(C₁-C₆ alkyl), arylS(O)₂(C₁-C₆ alkyl), heterocyclylS(O)₂(C₁-C₆ alkyl), aryl(C₁-C₆ alkyl)S(O)₂, heterocyclyl(C₁-C₆ alkyl)S(O)₂, C₂-C₆ alkenyl, C₁-C₆ alkoxy, carboxy-substituted C₁-C₆ alkoxy, C₁-C₆ haloalkoxy, C₁-C₆ hydroxyalkoxy, C₁-C₆ alkylcarboxy-substituted C₁-C₆ alkoxy, aryloxy, heterocyclyloxy, C₁-C₆ alkylthio, C₃-C₇ cycloalkyl(C₁-C₆ alkylthio), C₃-C₆ alkynylthio, C₁-C₆ alkylcarbonylamino, C₁-C₆ haloalkylcarbonylamino, SO₃H, -NR¹⁶R¹⁷, -C(O)NR²¹R²², S(O)₂NR¹³R¹⁴, S(O)₂R¹⁵, R²⁶C(O), carboxyl, C₁-C₆ alkoxycarbonyl, aryl and heterocyclyl; wherein the foregoing aryl and heterocyclyl moieties are optionally substituted by one or more of halogen, nitro, cyano, C₁-C₆ alkyl, C₁-C₆ haloalkyl, phenyl(C₁-C₆ alkyl), C₁-C₆ alkoxy, C₁-C₆ haloalkoxy, S(O)₂(C₁-C₆ alkyl), C(O)NH₂, carboxy or C₁-C₆ alkoxycarbonyl;
R⁷, R⁸, R⁹, R¹⁰, R¹¹, R¹³, R¹⁴, R¹⁶, R¹⁷, R¹⁸, R¹⁹, R²¹, R²², R²³ and R²⁴ are, independently hydrogen, C₁-C₆ alkyl, C₁-C₆ haloalkyl, C₁-C₆ hydroxyalkyl, C₃-C₇ cycloalkyl, C₃-C₇ cycloalkyl(C₁-C₄ alkyl) or phenyl(C₁-C₆ alkyl); and,
R¹⁵ and R²⁰ are, independently, C₁-C₆ alkyl, C₁-C₆ hydroxyalkyl, C₃-C₆ cycloalkyl, C₃-C₇ cycloalkyl(C₁-C₄ alkyl) or C₁-C₆ alkyl optionally substituted by phenyl;
R²⁵ and R²⁶ are, independently, C₁-C₆ alkyl or phenyl (optionally substituted by one or more of halogen, nitro, cyano, C₁-C₆ alkyl, C₁-C₆ haloalkyl, phenyl(C₁-C₆ alkyl), C₁-C₆ alkoxy, C₁-C₆ haloalkoxy, S(O)₂(C₁-C₆ alkyl), C(O)NH₂, carboxy or C₁-C₆ alkoxycarbonyl).